

Hotplate Emanation

The present invention relates to a method of deactivating dust mite allergens.

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Various allergens are known to trigger a human reaction. For example, it has been known for a long time that house dust can trigger allergenic reactions in humans, such as asthma and rhinitis. It was reported, as early as 1928
10 that it was the dust mites in the dust that were the primary source of the allergenic response, but it was only in the 1960's that researchers appreciated its significance.

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House dust mites produce detritus which causes allergenic reaction in many people. The major allergens are believed to be detritus from the mite species *Dermatophogoides farinae* and *Dermatophagoides pteronyssinus* (the allergens being known as Der f1 and Der p1 respectively). The
20 detritus includes faeces as well as body part residues of the mites. A review is given in *Experimental and Applied Acarology*, 10 (1991) p. 167-186.

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WO99/15208 describes a method for deactivating allergens derived from the *D. Pteronyssinus* and *D. Farinae* dust mite species, which comprises contacting the allergen with one of 28 deactivants which are described. These may be delivered into an airspace by aerosol spraying.

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WO 01/76371 describes further deactivants for house dust mite allergens. These may be delivered into an airspace by various methods including use of an ultra-sonic jet nebuliser, an ion wind generator, a candle incorporating a

deactivant, or heat to vaporise an oil. The examples given of the use of heat are to float the oil on water in an oil burner or to heat oil directly in an oil burner. Alternatively the volatile oil may be vaporized from a heated wick dipped into a reservoir of the volatile oil.

In accordance with a first aspect of the present invention there is provided a method of deactivating Der p and Der f allergens, the method comprising dispersing into an airspace an allergen-deactivating amount of an allergen-deactivating compound (hereinafter the "deactivant"), the deactivant being dispersed from a vessel in contact with a hotplate, the hotplate preferably being set to a temperature of at least 100°C.

Use of the noun deactivant and the verb deactivate in this specification denote that some or all of a source of allergens at a locus are rendered unable to evoke an allergenic response in a human, by a method of the present invention. The net result is that the source may be reduced in its allergenicity, or its allergenicity may be completely removed.

Many methods of killing dust mites are known. However it is thought that the allergenic response is due to dust mite detritus, not the dust mites themselves. Thus killing them does not remove the detritus. In the method of the present invention, the allergens Der p and/or Der f are deactivated. The killing of dust mites is not necessary.

The use of a hotplate enables the heat applied to disperse the deactivant to be controlled, in a manner which is not possible with prior methods.

- 5 Preferably the deactivant is dispersed into the airspace as a vapour.

Our work suggests that use of a hotplate below 100°C gives some allergen deactivating activity but that use of a
10 higher temperature gives allergen deactivating activity of a substantially and surprisingly higher level, even though the quantity of deactivant dispersed may be the same in each case.

- 15 Preferably the hotplate has an electrical heat source.

Preferably the vessel and the hotplate are in face-to-face contact. Preferably the hotplate has a flat surface and the vessel has a flat base, and the vessel rests on the
20 hotplate. Preferably the vessel has an opening in its upper region. Preferably it has a fully open upper face. Preferably, therefore, the vessel has a flat base, a side (if cylindrical) or sides depending upwardly therefrom, and no further side.

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Preferably the hotplate is set to a temperature of at least 130°C.

Preferably the hotplate is set to a temperature up to
30 300°C, preferably up to 250°C.

Preferably the deactivant is selected from:

a terpene hydrocarbon;
a citrus oil;
a mint oil;
bois de rose oil;
5 oil of jasmine;
frankincense;
oil of bergamot; and
oil of lemon grass.

10 Preferred terpene hydrocarbons include tea tree oil, pinol
and β -pinene.

An especially preferred deactivant is a citrus oil, most
preferably orange oil.

15 Another especially preferred deactivant is β -pinene.

A deactivant may suitably be a single compound.
Alternatively a mixture of deactivants may be used
20 together.

A deactivant may be part of a blend of compounds, not all
of which are deactivants. For example a citrus oil is a
blend of compounds not all of which will function as
25 deactivants.

A deactivant may suitably be dispersed into the airspace
over an extended period, for example at least 30 minutes,
and preferably at least 1 hour.

30 A deactivant may suitably be dispersed into the airspace
on two occasions, interrupted by a period in which there
is no deactivant dispersal. A deactivant may be dispersed

into the airspace on one or more further occasions, following a corresponding period or periods of no deactivant dispersal. Preferably each such dispersal occasion involves deactivant dispersal over an extended period, as described above. Preferably the or each period in which there is no deactivant dispersal is an extended period, for example at least 2 hours, preferably at least 4 hours, and most preferably at least 8 hours.

We have found that the method produces a prolonged reduction in the allergen loading of an allergen-contaminated inanimate substrate. Delivery of the deactivant into an airspace as described causes a permanent reduction in the population of allergens in an inanimate test source. By inanimate test source we mean a test source which is itself inanimate (e.g. it is not the skin or coat/fur of a live animal) and it does not contain living organisms, such as dust mites. Populations of dust mites would make any result difficult to interpret.

We have found that the reduction in allergen content in such a source is of long duration, for example at least 7 days, typically at least 14 days, and suitably at least 28 days. Indeed, in tests we have carried out over a 28-day period, we have found that the allergen content may continue to decline over time, even though the deactivant may have been used days or weeks before. The results suggest that the allergenic species have been truly denatured or degraded, to the extent that, firstly, they cannot re-form, and secondly, their degradation products are not themselves allergenic. It further suggests that the action of the deactivant is not merely a masking or damping effect. Any such effect would be likely to break down over time.

- The deactivant may be used as such, or may be comprised within an oil-on-water formulation, or may be comprised within an oil-in-water emulsion formulation. An oil/water formulation suitably comprises at least 0.5% by weight of the deactivant (in total, when more than one of said deactivants is employed), preferably at least 2%, more preferably at least 6%, most preferably at least 8%, and especially at least 10%. Suitably an oil-water formulation comprises up to 25% by weight of the deactivant (in total, when more than one of said deactivants is employed), more preferably up to 20% and most preferably up to 15%.
- 15 In this specification unless otherwise stated a percentage value given for a component denotes the weight of the component expressed as a percentage of the total weight of the emulsion.
- 20 The formation of emulsions is generally well known in the art and is described, for example, in Modern Aspects of Emulsion Science, edited by Bernard P. Binks, The Royal Society of Chemistry, 1998 and Surfactant Science and Technology, Second Edition, Drew Myers, 1992, VCH Publishers, Inc. Non-ionic surfactants may be especially
- 25 suitable. Proprietary surfactant packs may be employed to form emulsions, for example E-Z-MULSE (Trade Mark), a non-ionic surfactant pack from Florida Chemical Company, US.
- 30 The present invention involves the dispersal of an allergen deactivant into an airspace, preferably as a vapour. It is possible that airborne allergens may be deactivated but it is believed that there is effective

deactivation of allergens borne on surfaces within the airspace.

In accordance with a further aspect of the present invention there is provided the use of apparatus for deactivating Der p and/or Der f allergens at a locus, the apparatus comprising an allergen deactivant within a vessel, and a heat source used to accelerate the vaporization of the deactivant, the heat source being a hotplate in contact with the vessel.

In accordance with a further aspect there is provided an apparatus for deactivating Der p and/or Der f allergens comprising a vessel containing an allergen deactivant and a hot plate in contact with the vessel.

The present invention will be further described with reference to the following Examples.

20 Experimental Protocol

When using house dust for allergen denaturing tests an inherent difficulty is the variability of the amount of allergen in each small sample, even when taken from the same dust reservoir. The amount of dust in the pre-treatment sample must be accurately estimated in order to determine the extent of any allergen denaturing. In these tests the dust sample was applied to the test exposure surface and then one half of this surface dust was removed to measure the control pre-treatment allergen level of that specific sample. Each control was directly relevant to each sample, which gave the best possible estimate of the level of allergen in the sample before exposure to

possible denaturant. All tests employed a glass reinforced plastic booth of size 0.7m x 0.7m x 1.0m. Average values are stated.

- 5 The following Examples all measure the reduction of the house dust mite *Dermatophagoides pteronyssinus* allergen - Der p1.

10 House dust was passed through a number of sieves and the fraction smaller than 53 μ m was collected. 0.025g of dust was placed in a small sieve to distribute it evenly over the test surface. The test surface was a PTFE (polytetrafluoroethylene - trade mark TEFLON) coated metal tray of size 30cm x 30cm. The dust was applied to the
15 tray by moving the sieve continuously over the surface while tapping the sieve. One half of the dust was then removed by suction onto an in-line filter and the weight recorded, this was the pre-treatment control.

- 20 The tray was then placed in the booth. Three tea light holders - upwardly open cylindrical vessels (diameter 40mm, height 15mm) produced to hold nightlight candles - each containing 6ml of water and 0.8ml of orange oil - were placed together on an electric hotplate set to 250°C.
25 In practice we found that this meant that the hotplate temperature cycled between 130°C and 250°C.

Three more tea light holders each containing 6.8ml of 12% orange oil/water emulsified with E-Z-MULSE were placed
30 together on an electric hotplate set to 250°C. In practice we found that this meant that the hotplate temperature cycled between 130°C and 250°C. Heating was stopped after 105 minutes. These compositions did not

evaporate to dryness. This is believed to be due to remaining non-volatile surfactant from the E-Z-MULSE constituent).

- 5 Three more tea light holders each containing 6ml of water and 0.8ml of orange oil were placed on the hotplate, set to 90°C. In practice we found that this meant that the hotplate temperature cycled between 60°C and 90°C.
- 10 Three more tea light holders each containing 6ml of water and 0.8ml of orange oil were placed on the hotplate, set to 74°C. In practice we found that this meant that the temperature cycled between 40°C and 74°C.
- 15 Three more tea light holders each containing 6ml of water and 0.8ml of orange oil were placed on the hotplate, set to 50°C. In practice we found that this meant that the hotplate temperature cycled between 30°C and 50°C.
- 20 Three more tea light holders each containing 6ml of water and 0.8ml of orange oil were placed on the hotplate, set to 25°C. In practice we found that this meant the hotplate temperature cycled between 20°C and 25°C.
- 25 As a further comparison three tea lights containing only water (6.8ml each) were tested on the hotplate, set to 74°C. In practice we found that this meant the temperature cycled between 40°C and 74°C.
- 30 For each test the booth was sealed. Heat was delivered for sufficient time to allow the liquid to fully evaporate and then the hotplate was allowed to cool. After 24 hours the tray was removed, the dust was collected from it and

its weight recorded. The booth was washed with strong detergent between tests.

5 The test samples were assayed for the Der p1 allergen using an ELISA (Enzyme linked immunosorbent assay) to determine the allergen content. This was then related to the weight of dust that had been present in each sample. All of the samples were multiplied up to compare the amount of allergen expected to be present in a 0.1g sample
10 of dust. The percentage difference between the control sample and the exposed sample was then obtained.

The Der p1 allergen reductions were as follows:

15 Hot plate set to 130-250°C, 0.8ml orange oil on 6ml water - 75.4%
Hot plate set to 130-250°C, 0.8ml orange oil and 6ml water, emulsified - 91.0%
Hot plate set to 60-90°C, 0.8ml orange oil on 6ml
20 water - 48.6%
Hot plate set to 40-74°C, 0.8ml orange oil on 6ml water - 47.9%
Hot plate set to 30-50°C, 0.8ml orange oil on 6ml water - 48.0%
25 Hot plate set to 20-25°C, 0.8ml orange oil on 6ml water - 44.6%
Hot plate set to 40-74°C, 6.8ml water only - 42.7%

In the tests at 20-25°C there was incomplete evaporation.

30 Statistical analysis suggested that the finding of increased activity at a hotplate temperature of 130-250°C